

Centers for Disease Control and Prevention (CDC) National Center for Infectious Diseases 1324 Calle Cañada San Juan, Puerto Rico 00920-3860 Tel. (787) 706-2399 FAX (787) 706-2496

Criteria for the processing of dengue samples at the CDC Dengue Branch, San Juan, Puerto Rico

The diagnosis and treatment of dengue and dengue hemorrhagic fever are guided by the symptoms and findings that the patient presents, and cannot depend on laboratory confirmation, since routine tests can not confirm dengue with the speed required for patients in critical condition. The processing of samples for serologic diagnosis takes approximately one week, and the PCR results take approximately 3 days. Even so, it is necessary to eventually have a confirmation of the diagnosis, to exclude other etiologic possibilities, and to guide the follow-up of the patient's convalescence. The CDC Dengue Branch provides dengue testing free of cost to submitting physicians, state and private laboratories.

To obtain correct data on the spread of disease, adequately decide what laboratory tests to use, correctly interpret test results, and to assure that results get to the person who requested them, the following information is indispensable:

- Complete name, age and sex of the patient
- Home address
- Date of onset of symptoms
- Date that sample was obtained
- Complete name and mailing address of the physician, laboratory, clinic or hospital that the result should be sent to.

Samples without the above-mentioned information, or written with illegible handwriting or with more than a month from date of sample collection to date of arrival at CDC, will not be analyzed.

The clinical samples that are processed weekly are of great importance to determine the serotype and genetic make-up of viruses being transmitted in the region, and to determine their geographic distribution. This information can be used to detect risk factors that may lead to a

new and more severe pattern of disease, and to refine the clinical diagnostic ability of attending physicians. Diagnostic bleeding is not provided to persons referred to CDC.

In case of a severe dengue epidemic, CDC Dengue Branch will promptly analyze samples received with the minimum above-mentioned information. If number of specimens exceeds laboratory capacity, testing may be prioritized in the following order:

- fatal cases,
- cases in intensive care,
- hospitalized cases (with thrombocytopenia, hemorrhage, shock or hemoconcentration),
- all other cases.

We want to emphasize that to maintain efficient dengue surveillance, we will continue processing samples from outpatients and mildly ill persons any time of the year, and especially when dengue incidence in the island is relatively low (usually during the months from April to July).

Case Notification and Shipment of Samples of Suspected Dengue Cases

Dengue is characterized by an acute febrile picture accompanied by headache, retroorbital pain, body pain, often a rash, and other variable symptoms that can include obvious or mild hemorrhagic manifestations (such as a petechial rash) or hemoconcentration, shock or coma. This disease should be considered whenever there is an increase in the number of persons who go to the physician with an acute febrile illness, complaints of "monga" or "influenza," or an increase in the number of clinically diagnosed cases of German measles or regular measles.

Instructions for obtaining and handling samples:

- Once there is a clinical diagnosis of suspected dengue, take a blood sample (see #4 and #5) and fill out the Dengue Case Investigation Form (see copy attached).
 With this Form you comply with the legal reporting requirement. These forms can be obtained from the CDC Dengue Branch, the CDC Website http://www.cdc.gov/ncidod/dvbid/dengue/resources/DEN%20CASE%20Form%20Eng%202
 O04.pdf and also from the Regional Environmental Health Office of the Puerto Rico Department of Health. They can also be photocopied without restriction.
- 2. It is of great importance to fill out the Dengue Case Investigation Form in a clear and complete manner. The information received on each case (especially the date of onset of symptoms and date of sample collection) is crucial to select and interpret the laboratory analyses. Furthermore, a complete address makes it possible to identify the area where control measures should be implemented. Samples without the above-mentioned information, or written in illegible handwriting or with more than a month from date of collection to date of arrival at CDC, will not be analyzed.
- 3. The blood sample is taken in a red-top tube (preferably, but if not, you can use a green-top tube). Violet-top tubes (with heparin) should not be used. If dry ice is not available we recommend that after separating the serum, it must be maintained on ice or in a refrigerator until it is delivered to the CDC Dengue Branch. For samples from the USA or the exterior, we recommend to freeze the serum immediately after separated and to send in dry ice. The case investigation forms and the acute blood sample should reach CDC Dengue Branch as soon as possible. They can be sent through the local Environmental Health office. The acute sample can be sent immediately; there is no need to wait until the convalescent sample is taken.
- 4. To diagnose dengue, the laboratory requires a blood sample taken during the acute period of the disease and a second sample that can be taken from day 6 after the onset of symptoms. Informing the patient about the importance of coming back for a second sample, and giving an appointment for a specific day and hour, will increase the probability of obtaining the second sample. If the patient makes the first visit to the physician on or after day 6 after onset of the symptoms, that sample is enough. In that case, it is not necessary that the patient come for a second sample.
- 5. Acute-phase samples (taken on or before day 5 after onset of symptoms), will be used mainly for PCR analysis in order to detect virus. Convalescent-phase samples (taken on or after day 6 after beginning of symptoms) will be used mainly for detection of IgM anti-dengue antibodies by enzyme-linked immunosorbent assays (ELISA). Differential diagnosis for dengue and WNV

virus is available; but these tests need to be requested according to clinical presentation.

Interval since the

Type of sample onset of symptoms Type of Analysis

"Acute" until day 5 PCR
"Convalescent" 6 or more days Serology

Samples taken on days 4 and 5 of illness are of low yield for isolation as well as serology.

WHENEVER THERE IS A HOSPITALIZED SEVERE CASE, PLEASE INDICATE IT IN THE CASE INVESTIGATION FORM.

- 6. Reports will be sent to the physician (if the return address has been indicated) with the results of positive, or clearly negative, cases. In cases with negative virus isolation, we will await a convalescent-phase sample before reporting a result.
- 7. Results will be reported only to the laboratory or the physician who sent the sample (or an authorized secretary).

Dengue Branch, CDC March 10, 2008 1324 Calle Cañada

Puerto Nuevo Tel. (787) 706-2399 San Juan, PR 00920-3860 Fax (787) 706-2496

Points of contact:

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Dr. Hunsperger <u>enh4@cdc.gov</u>

CHECK LIST FOR OBTAINING AND SHIPPING DENGUE DIAGNOSTIC SAMPLES

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Type of sample	Interval since date	Type of		
	of onset of symptoms	Analysis		
Acute	up to 5 days	PCR		
Convalescent	6 or more days	Serology		

[] **Form -** "Dengue Case Investigation Form"

Can be obtained from the CDC Dengue Branch in San Juan or Internet: http://www.cdc.gov/ncidod/dvbid/dengue/resources/DEN%20CASE%20Form%20Eng%202004.pdf Please indicate on the sheet if the case is hospitalized. If it is a very severe case, indicate so on the "Comments" section.

Only the samples received with the information requested below, and written in a legible manner, will be analyzed:

- Complete name, age, and sex of patient
- Home address
- Date of onset of symptoms
- Date sample was obtained
- Complete name and mailing address of the physician, laboratory, clinic, or hospital

[]	Tube - Red or green top (not violet).	
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	Labeling -	Tube and	1 case	torm musi	t agree	(indicate	the san	ne name	of t	he ca	ase)	•

- [] **Volume** 2 cc. (ml.) of centrifuged serum or plasma
- [] **Storage** On ice or in a refrigerator (<u>not in a freezer</u>) until it is delivered to the CDC Dengue Branch.
- [] **Time of shipment** Not to exceed a month after taking the sample
- Way of shipment Check with local Department of Health.

Reasons for **REJECTING** samples:

- Samples without form, form without sample
- Incomplete or illegible form especially regarding date of onset of symptoms, date of sample collection
- Hemolyzed or frozen sample, or received more than a month after onset of illness

Dengue Branch, CDC March 28, 2008

1324 Calle Cañada

Puerto Nuevo Tel. (787) 706-2399 San Juan, PR 00920-3860 Fax (787) 706-2496



DENGUE CASE INVESTIGATION REPORT

CDC Dengue Branch and Puerto Rico Department of Health 1324 Calle Cañada, San Juan, P. R. 00920-3860 Tel. (787) 706-2399, Fax (787) 706-2496



Form Approved OMB No. 0920-0009

FOR CDC DENGUE BRANCH USE ONLY										
Case number Specimen # Days post onset (DPO) Type	Date Received	Specimen # Days post onset (DPO) Type	e Date Received							
SAN ID GCODE S1	_//	\$3	//							
SAN ID GCODE S2		S4	//							
Please compl	ete all sections									
Hospitalized: No Yes → Hospital Name:		Fatal:								
Name of Paliant		Yes _	No UNK							
Name of Patient: Last Name First Name Middle Name or Initial										
Mental Status Changes: If patient is a minor, name of father or primary caregiver: Yes No UN										
Last Name First Name Middle Name or Initial										
Home Address		Physician who referred the case								
City, Town: Barrio:	Name of Healthcare	Provider:								
Urbanization or sector:	Phone number:	Email address:								
Street: House / Apt. Number:	Send laboratory resu	ults to:								
Premise No.: Box: P.O. Box:										
Road No.: Km: Hm: Tel: Other Tel:										
Residence is close to: Zip Code:	Informo	ation about the person filling out this	s form							
Work address:	Name and title:	Phone number								
Work dudiess.										
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Day Month Year										
Day Month Year										
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Date of first symptom:/	2. Country of birth:	·	No UNK							
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Date of first symptom: Date specimen taken: Serum: First sample (Acute = first 5 days of illness - check for virus) Second sample (Convalescent = more than 5 days after onset - check for antibodies) Third sample Criteria for DENGUE HEMORRHA Second sample (Convalescent = more than 5 days after onset - check for antibodies) Third sample (Convalescent = more than 5 days after onset - check for antibodies) Third sample (Convalescent = more than 5 days after onset - check for antibodies) Criteria for DENGUE HEMORRHA ** Description of the properties of	2. Country of birth: 3. Have you been of the second of the	diagnosed with dengue before? Yes diagnosed with dengue before? Yes, another control of Yes, another control of Yes, another control of Yes, another city and Shock (#5) Symptoms continued Rash Chills Pallor or cool skin Nausea or vomiting Diarrhea Abdominal pain Cough Conjunctivitis (red eyes) Nasal Congestion Sore throat Jaundice Convulsion or coma	No UNK UNK Other cities or No UNK							

FOR CDC DENGUE BRANCH USE ONLY											
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This questionnaire is authorized by law (Public Health Service Act 42 USC 241). Although response to the questions asked is voluntary, cooperation of the patient is necessary for the study and control of the disease. Public reporting burden for the collection of information is estimated to average 15 minutes per response. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing the burden to PHS Reports Clearance Officer; Rm. 721-H, Humphrey Bg; 200 Independence Ave., SW; Washington, DC 20201; ATTN: PRA, and to the Office of information and Regulatory Affaire, Office of Management and Budget, Washington, DC.